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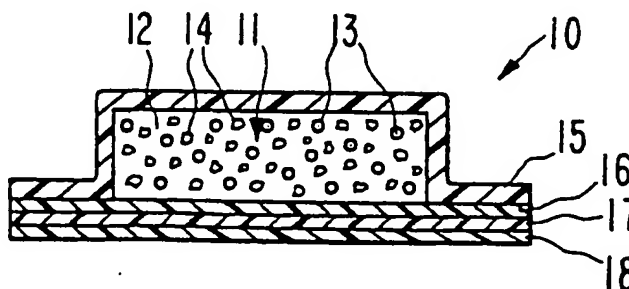
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(54) Title: DOSAGE FORM HAVING REDUCED ABUSE POTENTIAL



(57) Abstract

Compositions and dosage forms for administering abusable substances which have a reduced potential for abuse without diminishing the therapeutic or beneficial effects of the abusable substance. Topical compositions (1) contain: (a) the abusable substance (3) in a form which is permeable to the skin or mucosa; and (b) an antagonist (4) for the abusable substance in a form that is impermeable to the skin or mucosa to which the composition is to be applied. Controlled release dosage forms (10) which release the abusable substance (13) from a drug reservoir composition (11) confined behind a release rate controlling barrier (16) contain the abusable substance (13) in a form that is releasable from the dosage form (10) and the antagonist (14) in a form that is not releasable from the dosage form. This invention permits an abusable substance to be administered by the prescribed route at a substantially full potency, while reducing the abuse potential of the compositions and dosage forms via other routes of administration. Preferred transdermal forms of this invention utilize fentanyl as the abusable substance and naltrexone as the antagonist.

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DOSAGE FORM HAVING REDUCED ABUSE POTENTIAL

FIELD OF THE INVENTION

5 This invention relates to dosage forms of abusable substances having reduced potential for abuse. In particular, the dosage forms of this invention are intended to administer the abusable substance to the body by topical application to the skin or mucosa or to release the abusable substance to the body through a
10 membrane on the dosage form.

BACKGROUND OF THE INVENTION

 The potential for abuse by either oral or parenteral routes of narcotic and other psychoactive drugs is well known. For example,
15 the potential for abuse of the synthetic narcotic drug fentanyl is so high that it has become a major cause of death for anesthesiologists and other hospital workers having access to the drug. In order to prevent abuse of these substances, it has been proposed to provide dosage forms which combine the abusable substance with an amount of
20 an antagonist for the abusable substance sufficient to eliminate the "high" associated with abuse of the substance without eliminating the other therapeutic benefits for which the drugs are intended to be administered. See, for example, U.S. Patents 4,457,933, 3,493,657 and 3,773,955 which are incorporated herein by reference.

25 Many abusable substances are capable of being administered to the body by direct application of the drug to the skin or mucosa, i.e., nasal, vaginal, oral, or rectal mucosa. They can also be delivered to the body from advanced dosage forms such as those described in U.S. Patents 4,665,766, 4,588,580, 3,993,073, and
30 3,845,770 which are incorporated herein by reference. Compositions suitable for topical application to the skin or for forming reservoir compositions in the advanced dosage forms described above could be subject to abuse, and it would clearly be desirable to have such compositions or dosage forms available in a condition in which the
35 abuse potential of the composition or dosage form is reduced without diminishing the therapeutic efficacy of the abusable substance to be administered.

It is, accordingly, an object of this invention to provide a composition of matter for topical application of an abusable substance to the body, which composition has a low potential for abuse.

5 It is another object of this invention to provide a dosage form for the therapeutic administration of an abusable substance, which dosage form has a low potential for abuse.

It is another object of this invention to provide a transdermal dosage form of an abusable substance having a low
10 potential for abuse.

It is a further object of this invention to provide methods for administering abusable substances to the body from compositions and devices having a low potential for abuse.

15

BRIEF DESCRIPTION OF THE INVENTION

The abuse potential that this invention is intended to reduce is not the abuse potential associated with the illicit, nonprescription or recreational use of the topical composition or dosage form of this invention in the same mode of administration as
20 intended for its therapeutic use. Instead, the abuse potential referred to herein relates to the use of the topical or drug reservoir compositions of this invention by other modes of administration such as the injection or ingestion of a topical composition intended to be applied directly to the skin or the
25 removal of the drug reservoir composition from the dosage form and the injection or ingestion of the drug reservoir composition..

In its broadest aspect, this invention contemplates a composition of matter adapted for topical administration to the skin or mucosa which comprises a mixture, typically in a pharmaceutically
30 acceptable carrier, of an abusable substance in combination with an amount of antagonist for said abusable substance sufficient to substantially negate the pharmacological effect of the abusable substance; the abusable substance being present in said composition in a form which is permeable through the skin or other body membrane
35 to which it is to be applied and the antagonist being present in the composition in a form which is substantially impermeable to the skin

or other body membrane to which the topical composition is to be applied.

In another aspect of this invention, a composition containing the abusable substance and an amount of an antagonist therefor sufficient to substantially negate the pharmacological effect of the abusable substance forms the drug reservoir composition of a dosage form in which the drug reservoir composition is confined within and separated from the body surface or fluid to which the abusable substance is to be administered by a barrier through which the abusable substance is released and the antagonist is not released.

DESCRIPTION OF THE DRAWINGS

Fig. 1 is a schematic cross-section of one embodiment of this invention;

Fig. 2 is a schematic cross-section of another embodiment of this invention;

Fig. 3 is a schematic cross-section of another embodiment of this invention; and

Fig. 4 is a schematic cross-section of another embodiment of this invention.

DESCRIPTION OF THE INVENTION
INCLUDING THE BEST MODE THEREOF

The dosage forms of this invention may comprise an osmotically driven dosage form such as described in U.S. patents 5 4,655,766 of 3,845,770, the abusable substance being present in the drug reservoir in a soluble, osmotically active form and the antagonist being present in an insoluble, osmotically inactive form. The dosage forms of this invention can also operate on principles of diffusion, such as shown in U. S. patents 3,993,073 and 4,588,580, 10 with the abusable substance being present in the drug reservoir in a form that would be permeable to the material forming the barrier and the antagonist being present in a form that would be impermeable to the material forming the barrier.

Drugs having a potential for abuse include natural and 15 synthetic narcotics and other psychoactive substances. Representative of such substances are, without limitation, analgetic agents such as fentanyl, sufentanil, carfentanil, lofentanil, alfentanil, hydromorphone, oxycodone, propoxyphene, pentazocine, methadone, tilidine, butorphanol, buprenorphine, levorphanol, 20 codeine, oxymorphone, meperidine, dihydrocodeinone and cocaine.

As used herein, an antagonist for an abusable substance is a compound or composition which acts on the recipient to prevent or substantially diminish the pharmacological effects of the abusable substance or to substantially delay their manifestation. Thus, with 25 respect to natural and synthetic narcotic agents, typical antagonists include, without limitation, naloxone, naltrexone, nalbuphine, nalorphine and levelorphan.

The amount of antagonist required to prevent the occurrence of the pharmacological effect of an abusable substance will vary from 30 substance to substance and antagonist to antagonist but can be readily determined from information available to the art regarding the pharmacokinetics and pharmacodynamics of the particular abusable substance and the particular antagonist. In the event of abuse of the unused reservoir compositions, in those circumstances in which 35 the abusable substance has a significantly longer half-life in the body than the antagonist, the antagonist will prevent the occurrence

of the pharmacological effect for a substantial period of time but may thereafter allow some pharmacological effect to be manifested.

This is because pharmacologically effective amounts of the abusable substance may linger in the body after the levels of the antagonist have been reduced below the effective level. In this circumstance the abuse potential of the dosage form will still be reduced because of the substantial delay between the ingestion or injection of the composition of this invention and the onset of the abuse inducing pharmacological effects. If the substance being abused does not promptly produce the effects for which it is being abused, its abuse potential will obviously be substantially reduced. Delays in the onset of the abuse inducing effects of as little as 15 minutes to 1/2 hour can significantly reduce the abuse potential, and compositions according to this invention can easily produce delays significantly longer than that.

Certain antagonists, such as naltrexone, are effective both orally and parenterally although the parenteral half-life may be much shorter than the oral half-life. Other antagonists, such as naloxone, are much more effective parenterally than orally.

Also, most narcotics are significantly more potent when administered parenterally than orally. Fentanyl, for example, is only about 1/10 as potent on oral, as compared to parenteral administration. Thus, oral abuse is not a significant problem with fentanyl, and an antagonist such as naloxone that is primarily parenterally active can be used. If, however, an abusable substance is both orally and parenterally active, an antagonist such as naltrexone would be preferred or a mixture of orally active and parenterally active antagonists could be used according to this invention. Further, intravenous abuse generally produces the desired abuse inducing effect more rapidly and with greater intensity than oral abuse and is, for many abusers, the preferred route of administration of an abusable substance. Thus, the use of an orally inactive antagonist can still substantially reduce the potential for abuse of the compositions and dosage forms of this invention.

Accordingly, as used herein, an antagonist is present in the compositions or dosage forms of this invention in an "abuse

negating amount" if the amount of antagonist is sufficient, when administered by at least one manner of potential abuse other than the manner by which the composition or dosage form is intended to be administered, to prevent or diminish the occurrence of the pharmacological effects of the abusable substance or to significantly delay the onset of these effects.

As used herein, abusable substances and antagonists are considered to be permeable through the skin or mucosa if present in the composition or dosage form as a chemical entity which is capable of producing its intended pharmacological effect when topically applied to the skin or mucosa either by direct application of the composition to the skin or by application of a dosage form containing the composition. Similarly, abusable substances and antagonists are considered to be impermeable through skin or mucosa if present in the dosage form as a chemical entity which is incapable of producing its intended pharmacological effect when topically applied at the administration site.

For example, as pointed out in patent 4,588,580, fentanyl and its derivatives are permeable through the skin as the base form of the drug and are impermeable through the skin in ionic form such as the commercially available fentanyl citrate, for example. Antagonists such as naloxone, naltrexone and nalbuphine, as shown in U.S. Patent 4,573,995 which is incorporated herein by reference, are impermeable to the skin in the absence of an appropriate permeation enhancer.

Similarly, abusable substances and antagonists are considered to be permeable to the barrier forming a dosage form if the rate of permeation of the abusable substance or antagonist through the barrier is sufficient to permit the abusable substance or the antagonist to produce its desired systemic effect from the site of administration and impermeable to the barrier if the rate of permeation is insufficient to permit the abusable substance or antagonist to produce its desired effect at the site of administration.

Referring now to Figure 1, a composition 1 for topical application to the skin or mucosa according to this invention

comprises a pharmaceutically acceptable carrier 2 having uniformly dispersed therethrough an amount of an abusable substance 3 in a skin or mucosa permeable form and an abuse negating amount of an antagonist 4 for said abusable substance in a skin or mucosa impermeable form. The composition 1 may be applied directly onto the skin or mucosa from a container for the same, such as a bottle or tube, and subsequently covered, if desired, with a protective overlay. It is preferable, however, to quantify the dose and the area of application by placing the composition in an impermeable container 5 of the correct size to provide a unit dose which may be held on the skin by adhesive means 6 or other appropriate fastening means.

In operation the composition of Figure 1 would administer the abusable substance through the skin or mucosa to produce the intended therapeutic effect, and the antagonist would not significantly affect or diminish the therapeutic effect of the abusable substance. If, on the other hand, an attempt were made to abuse the composition 1 by administration through some other portal, such as orally or intravenously, the antagonist would produce its antagonistic effect and prevent the occurrence of the abuse inducing effect.

In addition to the abusable substance and the antagonist, composition 1 may also contain a permeation enhancer for the abusable substance which should not significantly enhance the permeability of the antagonist, thickeners and other additives, all as known to the art. The composition 1 can be in the form of an ointment, cream, gel, paste, solution or lotion, for example.

Referring now to Figure 2, a dosage form 10 for administering an abusable substance to the skin or mucosa according to this invention is shown in which the reservoir composition has a reduced potential for abuse. The dosage form 10 comprises a drug reservoir composition 11 typically in the form of a gel or polymeric carrier 12 having uniformly dispersed therethrough an abusable therapeutic agent 13 and an abuse negating amount of an antagonist therefor 14. The composition 11 is preferably disposed between an impermeable backing 15, an abusable substance releasing means such as

release rate controlling membrane 16 and an abusable substance permeable adhesive 17, all as is known to the art. An abusable substance and antagonist impermeable release liner 18 is applied to the adhesive layer and is removed prior to use. Suitable materials for use in manufacturing the various layers are described in the above-identified patents.

In this embodiment of the invention the abusable substance 13 is present in the reservoir composition in a form which is permeable through the rate controlling membrane 16. The antagonist 14 is present in a form which is not permeable through either the rate controlling membrane 16, the adhesive 17 or the skin.

In operation, the dosage form 10 would administer the abusable substance through the skin at the rate intended for therapeutic effect, and the antagonist would not significantly affect or diminish the therapeutic effect of the abusable substance. If, on the other hand, an attempt were made to abuse dosage form 10 by administration of drug reservoir composition 11 through some other portal, such as orally or intravenously, the antagonist would produce its antagonistic effect and prevent the occurrence of the abuse inducing effect.

Referring now to Figure 3, a dosage form 20 is shown for orally administering an orally active abusable substance. Dosage form 20 comprises a drug reservoir composition 21 comprising a carrier having uniformly dispersed therethrough an abusable substance 22 and an abuse negating amount of at least one antagonist 23 for said abusable substance, said composition being enclosed within abusable substance releasing means 24. The abusable substance is present in the composition in a form that is permeable through means 24, and the antagonist is present in a form that is impermeable to means 24. If it is desired to prevent both oral and parenteral abuse of the reservoir composition, antagonist 23 should be either an orally or parenterally active antagonist or a mixture of orally and parenterally active antagonists.

In operation, the dosage form 20 would be administered orally. The abusable substance would be delivered through the releasing means 24 to the gastrointestinal tract gradually at the

controlled rate intended for therapeutic effect and the antagonist would not significantly affect or diminish the therapeutic effect of the abusable substance. If, on the other hand, an attempt were made to abuse the composition 21, in order to rapidly obtain the abuse inducing effect, by removing it from the dosage form 20 and ingesting the composition or administering it through some other portal, such as intravenously, the antagonist would produce its antagonistic effect and prevent the occurrence of the abuse inducing effect.

Referring now to Figure 4, another embodiment of this invention is shown in which the dosage form 30 operates on osmotic principals. In this embodiment, the dosage form 30 comprises a drug reservoir composition 31 comprising an osmotically active abusable substance 32 dispersed within an abuse negating amount of at least one osmotically inactive antagonist 33 for said abusable substance enclosed within a semipermeable housing 34. Housing 34 is permeable to water, impermeable to the abusable substance and the antagonist and is provided with an outlet 35 through which the abusable substance solution formed within the housing can be osmotically discharged. The abusable substance is present in the drug reservoir composition in a soluble, osmotically active form and the antagonist is present in an insoluble, osmotically inactive form.

In operation, the dosage form 30 would be orally administered. When the dosage form is in contact with body fluids, water will pass through the semipermeable housing 34 by osmosis, forming a solution of the abusable substance 32 which will be pumped out of the outlet 35 at the same rate that water passes into the dosage form 30. The antagonist, being insoluble, will not enter into solution and will remain within housing 34. Thus, the abusable substance will be administered at the rate intended for therapeutic effect, and the antagonist would not significantly affect or diminish the therapeutic effect of the abusable substance. If, on the other hand, an attempt were made to abuse the composition 31 by removing it from the dosage form 30 and ingesting the composition or administering it through some other portal, such as intravenously, the antagonist would produce its antagonistic effect and prevent the occurrence of the abuse inducing effect.

Having thus generally described our invention, the following examples illustrate various embodiments of our invention.

Example 1

Aqueous gels for topical application to the skin to administer the abusable substance, fentanyl, transdermally are prepared from a mixture having the following formulation:

Component	% by weight
Ethanol	22.1
Hydroxyethyl	
Cellulose	1.9
Fentanyl base	1.0
Naloxone base	1.0
or	
Naltrexone HCl	20.0
Water	Balance

The gels of this example contain abuse negating amounts of the antagonists and can be topically applied to the skin to administer fentanyl to the body without administering any therapeutically significant amount of the antagonists. Ethanol is a selective skin permeation enhancer in that it significantly increases the flux of fentanyl through skin without producing any therapeutically significant effect on the skin flux of the antagonists. The parenteral half-lives of naloxone and naltrexone are significantly shorter than the half-life of fentanyl, and naloxone and fentanyl are relatively inactive on oral administration. Thus, the gels containing naloxone or naltrexone will significantly delay the onset of the abuse inducing effects of the fentanyl in the event of I.V. abuse of the compositions of Example 1. The gels containing naltrexone HCl will also be more effective than the gel containing naloxone in preventing the oral abuse of the compositions of Example 1.

Example 2

Transdermal delivery devices having a drug releasing area of approximately 10 cm² are prepared by pouching, in a rotary heat sealing machine, the compositions of Example 1 between an impermeable backing formed from an aluminized polyester/ethylene vinyl acetate copolymer (EVA) film (Scotchpak 1018 available from 3M Corporation) and a multilaminate film comprising the rate controlling membrane, 2 mil EVA (9% VA), 1.8 mil of an amine resistant silicone adhesive (Dow

Corning X7-2920) and a fluorocarbon coated polyethylene terephthalate release liner (Scotchpak 1022) at a gel loading of approximately 15 mg/cm². The steady state release rate of fentanyl, naloxone and naltrexone from such systems into an aqueous bath as determined at 5 32°C. will be approximately as follows:

Fentanyl	4.0 mcg/cm ² /hr
Naloxone base	0.15 mcg/cm ² /hr
Naltrexone HCl	0.2 mcg/cm ² /hr

The rate of release of the fentanyl is sufficient to
10 produce therapeutic effects upon application of the dosage forms to the body and maintenance of the devices in place for 24 hours, for example. The rates of release of the antagonists, however, are insufficient to produce any significant reduction of the beneficial effect of the fentanyl when administered transdermally. Even if the
15 antagonists were released from the dosage form at significantly higher rates, they would not be antagonistically effective because neither is permeable to the skin in the absence of a permeation enhancer. Ethanol is not a permeation enhancer for either naloxone or naltrexone HCl. However, as with the compositions of Example 1,
20 the unused and used reservoir compositions will have reduced potential for abuse should the reservoir compositions be removed from the device and used in an abusive manner as described above.

Although this invention has been described with respect to various embodiments thereof it should not be construed as being
25 limited thereto. Various modification can be made to the subject matter disclosed herein within departing from the scope of this invention which is limited only by the following claims wherein:

WE CLAIM:

1. A composition of matter for the administration of an
abusable substance by application of the composition to the
skin or mucosa, said composition having a reduced potential for
abuse and comprising a mixture of:

a) an amount of an abusable substance in a form that
is permeable to the skin or mucosa to which it is intended to
be applied; and

b) an abuse negating amount of at least one
antagonist for said abusable substance in a form that is
impermeable to the skin or mucosa to which it is to be applied.

2. The composition of claim 1 further comprising a
pharmaceutically acceptable carrier within which said abusable
substance and said antagonist are dispersed.

3. The composition of claim 1 wherein said antagonist is
selected from the group of impermeable forms of naloxone,
naltrexone, nalbuphine and levorphan.

4. The composition of claim 1 wherein said antagonist is
naltrexone.

5. A dosage form for administering an abusable substance
to the body, said dosage form having a reduced potential for
abuse and comprising, in combination:

a) abusable substance releasing means through which
said abusable substance is released to the body; and

b) a drug reservoir separated from the body by said
releasing means, said reservoir comprising a mixture of said
abusable substance and an abuse negating amount of at least one
antagonist for said abusable substance:

i) said abusable substance being present in
said reservoir in a form that is releasable to the body through
said releasing means; and

ii) said antagonist being present in said
reservoir in a form that is not releasable through said
releasing means.

6. The dosage form of claim 5 wherein said releasing
2 means comprises a semipermeable material permeable to water and
impermeable to said abusable substance and said antagonist and
4 said drug reservoir comprises a water soluble form of said
abusable substance and a water insoluble form of said
6 antagonist and said semipermeable material is provided with
outlet means for a solution of said abusable substance.

7. The dosage form of claim 5 wherein said releasing
means is permeable to said abusable substance and impermeable
to said antagonist.

8. A dosage form for the administration of an abusable
2 substance through the skin or mucosa, said dosage form having a
reduced potential for abuse and comprising:

4 a) abusable substance releasing means through which
said abusable substance is released to the skin or mucosa;

6 b) a drug reservoir separated from the skin or
mucosa by said releasing means, said reservoir comprising a
8 mixture of said abusable substance and an abuse negating amount
of at least one antagonist for said abusable substance:

10 i) said abusable substance being present in
said reservoir in a form that is releasable to the skin through
12 said releasing means; and

14 ii) said antagonist being present in said
reservoir in a form that is not releasable through said
releasing means; and

16 c) means for maintaining said reservoir in abusable
substance releasing relationship to the skin or mucosa.

9. The dosage form of claim 8 wherein said releasing
2 means comprises a semipermeable material permeable to water and
impermeable to said abusable substance and said antagonist and
4 said drug reservoir comprises a water soluble form of said
abusable substance and a water insoluble form of said
6 antagonist and said semipermeable material is provided with
outlet means for a solution of said abusable substance.

2 10. The dosage form of claim 8 wherein said releasing
means is permeable to said abusable substance and impermeable
to said antagonist.

2 11. A method for administering an abusable substance to
the skin or mucosa to produce a predetermined beneficial effect
which comprises:

4 a) contacting the skin or mucosa with a composition
comprising a mixture of:

6 i) an amount of an abusable substance in a form
that is permeable to the skin or mucosa to which it is intended
8 to be applied; and

10 ii) an abuse negating amount of at least one
antagonist for said abusable substance in a form that is
impermeable to the skin or mucosa to which it is to be applied;
12 and

14 b) maintaining said composition on the skin or
mucosa for a period of time at least sufficient to produce the
beneficial effect.

2 12. A method for administering an abusable substance to
the body from a dosage form having a reduced potential for
abuse comprising:

4 a) placing a dosage form containing an amount of an
abusable substance and an abuse negating amount of at least one
6 antagonist for said abusable substance in abusable substance
delivering relationship to the body;

8 b) delivering said abusable substance to the body
from said dosage form; and

10 c) retaining said antagonist within said dosage form.

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FIG. 1

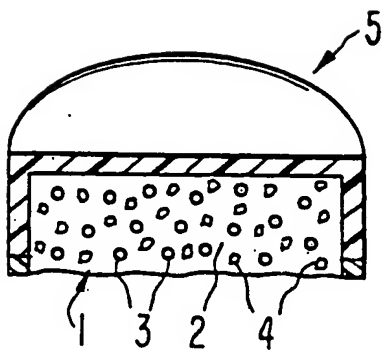


FIG. 2

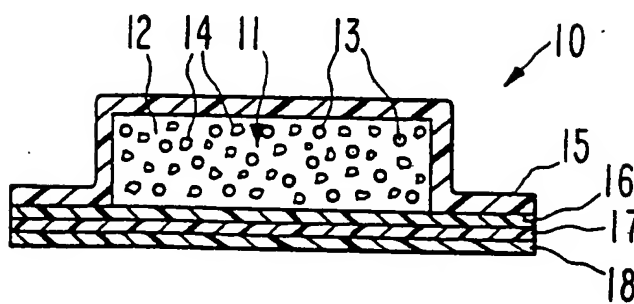


FIG. 3

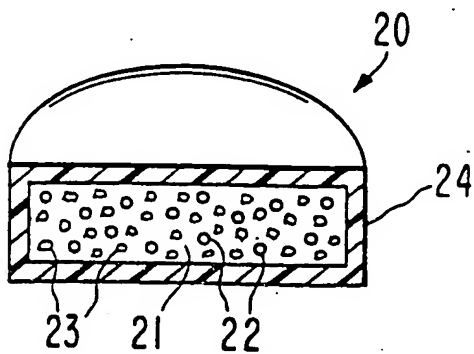
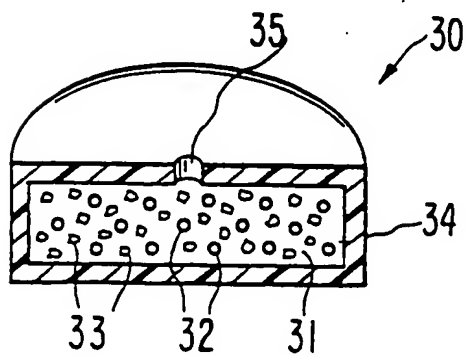


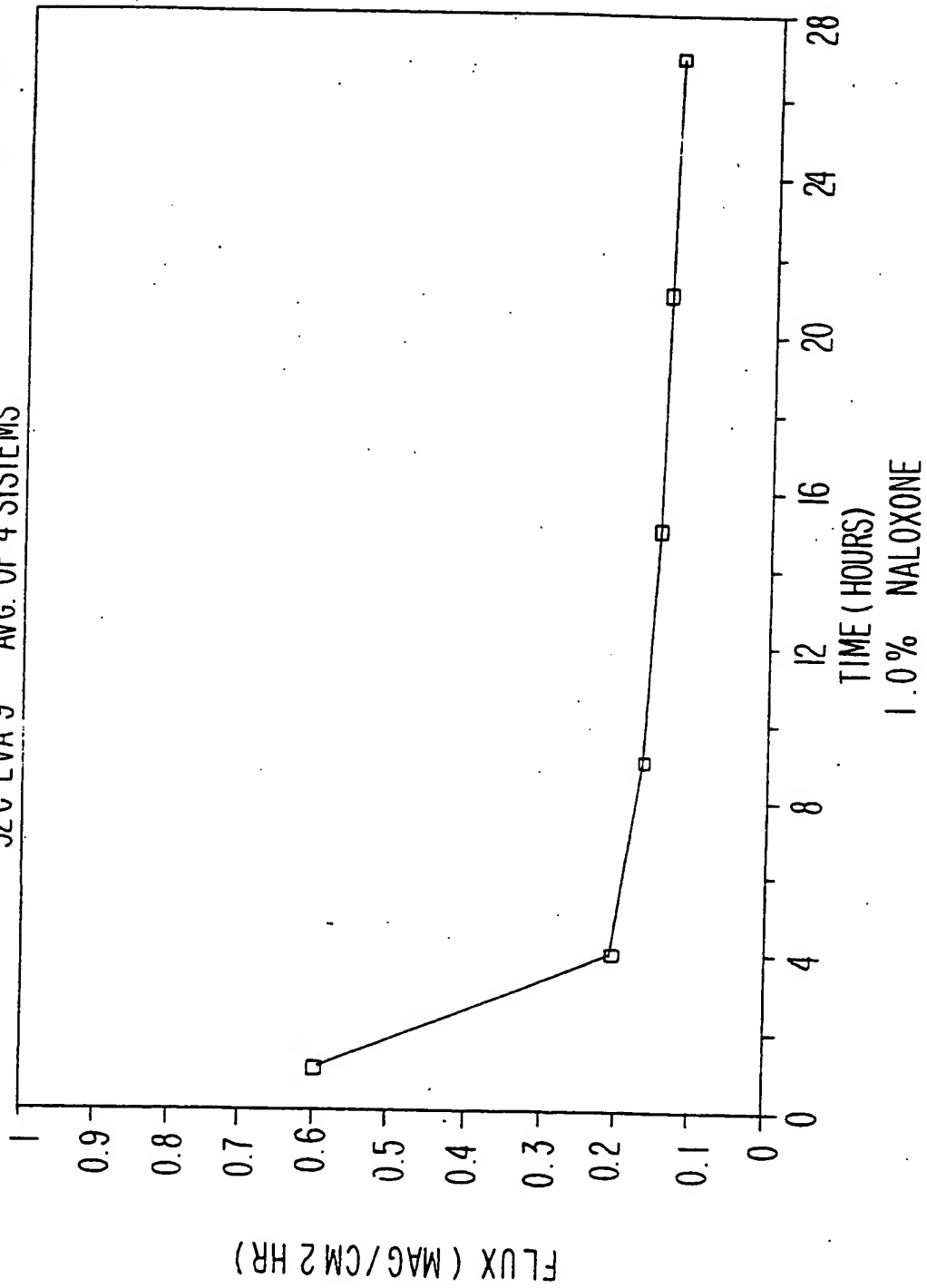
FIG. 4



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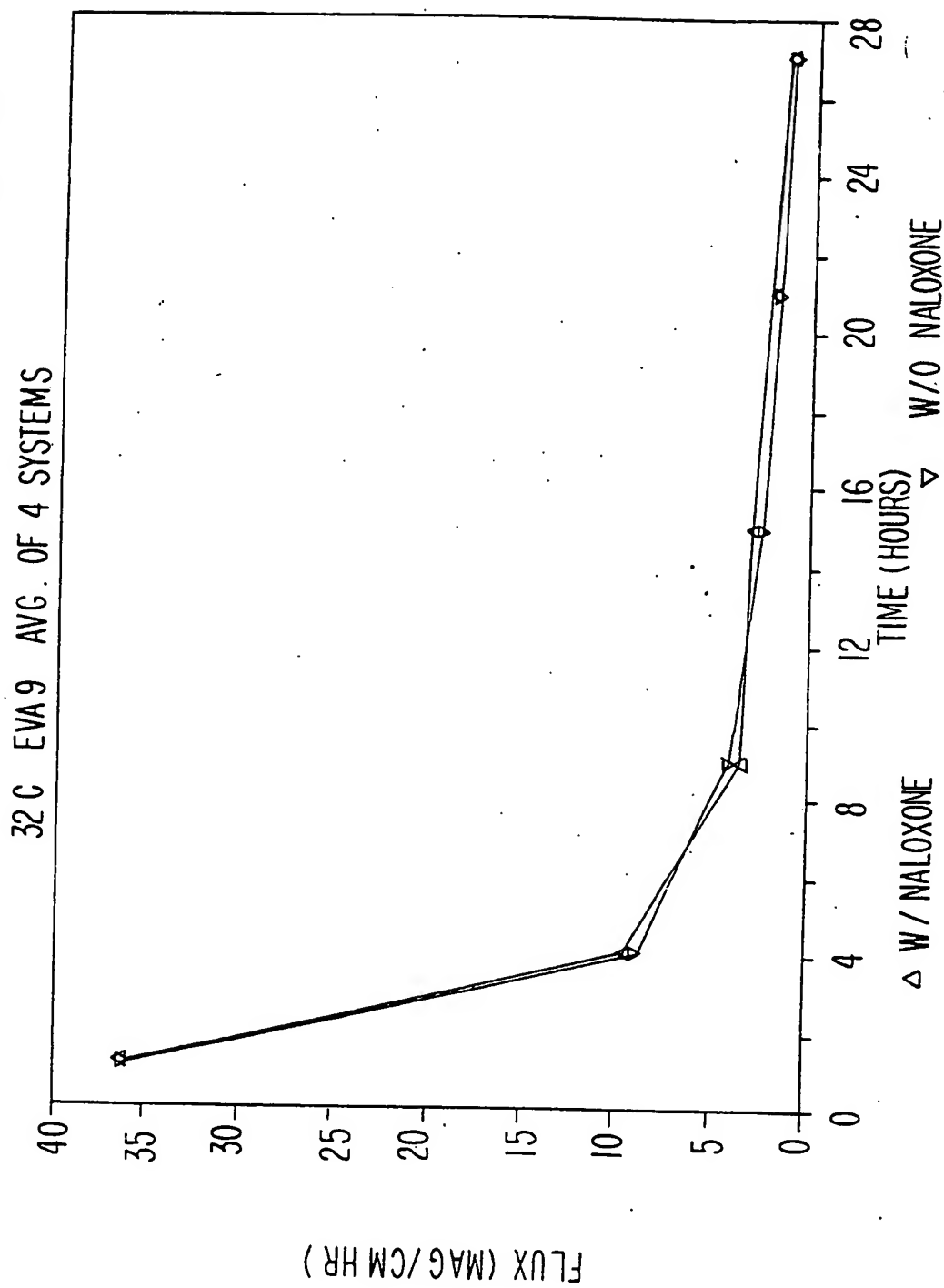
FIG. 5

32C EVA 9 AVG. OF 4 SYSTEMS



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FIG. 6



INTERNATIONAL SEARCH REPORT

International Application No PCT/US 89/04722

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) *		
According to International Patent Classification (IPC) or to both National Classification and IPC IPC5: A 61 K 31/485, 9/00, A 61 L 15/14		
II. FIELDS SEARCHED		
Minimum Documentation Searched ?		
Classification System	Classification Symbols	
IPC5	A 61 L; A 61 K	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched *		
III. DOCUMENTS CONSIDERED TO BE RELEVANT †		
Category *	Citation of Document, †† with Indication, where appropriate, of the relevant passages ‡	Relevant to Claim No. ‡
X	EP, A1, 0185472 (RECKITT AND COLMAN PRODUCTS LIMITED) 25 June 1986, see the abstract and the claims --	1-3
X	EP, A1, 0144243 (RECKITT AND COLMAN PRODUCTS LIMITED) 12 June 1985, see the abstract and the claims --	1-3
X	US, A, 4582835 (JOHN W. LEWIS ET AL.) 15 April 1986, see the abstract and the claims	1-3
Y	--	5-12
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>* Special categories of cited documents: ††</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"A" document member of the same patent family</p> </div> </div>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
29th January 1990	- 1. 03. 90	
International Searching Authority	Signature of Authorized Officer	
EUROPEAN PATENT OFFICE	T.K. WILLIS	

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
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**ANNEX TO THE INTERNATIONAL SEARCH REPORT
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This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office (EPO) file on 08/11/89. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

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